Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to: Laura Oliven, Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, D.C. 20503.

Dated: August 17, 1998.

Jane Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 98–22575 Filed 8–21–98; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Council; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), announcement is made of the following National Advisory body scheduled to meet during the month of September 1998.

Name: National Advisory Council on the National Health Service Corps (NHSC).

Date and Time: September 9, 1998; 6:00 p.m.–9:00 p.m.; September 10–12, 1998; 9:00 a.m.–5:00 p.m.; September 13, 1998; 8:00 a.m.–11:00 a.m.

Place: Sheraton National Hotel, 900 South Orme Street, Arlington, VA 22204, (703) 521–1900.

The meeting is open to the public.

Agenda: Items will include, but not be limited to: In preparation for the year 2000 reauthorization the National Advisory Council has developed a draft position paper, ''The National Health Service Corps for the 21st Century.'' Reactions, suggestions and criticisms to this paper will be heard from public and private partners and other interested organizations on September 10–12. Copies of the draft paper will be available at the meeting. Other agenda items include updates on the NHSC program.

For further information, call Ms. Eve Morrow at (301) 594–4144.

Agenda items are subject to change as priorities dictate.

Dated: August 17, 1998.

Jane M. Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 98–22574 Filed 8–21–98; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Council; Notice of Meeting

In accordance with section 10(a) (2) of the Federal Advisory Committee Act (Pub. L. 92–463), announcement is made of the following National Advisory body scheduled to meet during the month of September 1998.

Name: National Advisory Committee on Rural Health.

Date and Time: September 13, 1998; 5:00 p.m.–6:30 p.m.; September 14–15, 1998; 8:30 a.m.–5:00 p.m.; September 16, 1998; 8:30 p.m.–11:30 a.m.

Place: Holiday Inn, Georgetown, 2101 Wisconsin Avenue, Washington, DC 20007, Phone: (202) 338–4600, FAX: (202) 333–6113.

The meeting is open to the public.

Agenda: A special session will be conducted on Sunday, September 13, for orientation of new members who were just appointed. Monday will include a panel discussion of ''Rural Researchers' Access to National Health Survey Data,'' a presentation and discussion of the new guidelines for designating HPSAs, and a report on ''Critical Access Hospitals.'' Tuesday will include legislative, telehealth, and regulatory updates. A presentation and discussion on the ''National Bipartisan Commission on the Future of Medicare'' will be followed by a discussion of Department interests and priorities for FY 1999. Agenda items are subject to change as priorities dictate.

Anyone requiring information regarding the subject Committee should contact Ms. Arlene A. Granderson, Office, or Rural Health Policy, Health Resources and Services Administration, Room 9–05, Parklawn Building, Rockville, Maryland 20857; telephone (301) 443–0835, FAX (301) 443–2803. Persons interested in attending any portion of the meeting or having questions regarding the meeting should contact Ms. Arlene Granderson or Ms. Lilly Smetana, Office of Rural Health Policy, Health Resources and Services Administration, telephone (301) 443–0835.

Dated: August 17, 1998.

Jane M. Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 98–22576 Filed 8–21–98; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of Inspector General

Publication of OIG Compliance Program Guidance for Clinical Laboratories

AGENCY: Office of Inspector General (OIG), HHS.

ACTION: Notice.

SUMMARY: This **Federal Register** notice sets forth the OIG's recently-issued Compliance Program Guidance for Clinical Laboratories. The OIG had previously developed and published a model compliance plan for the clinical laboratory industry on March 3, 1997. This Compliance Program Guidance for Clinical Laboratories is intended to be more consistent with compliance program guidances issued by the OIG with respect to the hospital industry and to home health agencies, and serves to clarify various aspects of the original model plan. As with previously-issued compliance program guidances, we believe that the development of this guidance for clinical laboratories will continue as a positive step towards promoting a higher level of ethical and lawful conduct throughout the entire health care community.

FOR FURTHER INFORMATION CONTACT: Christine Saxonis, Office of Counsel to the Inspector General, (202) 619–2078.

SUPPLEMENTARY INFORMATION: As part of a major initiative to engage the private health care community in combating fraud and abuse, the OIG developed and published in the **Federal Register** a model compliance plan for the clinical laboratories (62 FR 9435; March 3, 1997). The compliance plan was intended to provide clear guidance to that aspect of the clinical laboratory industry that was interested in reducing fraud and abuse within their organizations. Since that issuance, the OIG has developed and issued specific compliance program guidance for the hospital industry and for home health agencies.

This compliance program guidance is intended to refine and build on the original model guidance plan for clinical laboratories. In developing an effective compliance program, the OIG has identified 7 fundamental elements. They are:

- Implementing written policies, procedures and standards of conduct;
- Designing a compliance officer and compliance committee;
- Conducting effective training and education;
- Developing effective lines of communication;
- Enforcing standards through well-publicized disciplinary guidelines;
- Conducting internal monitoring and auditing; and
- Responding promptly to detected offenses and developing corrective action.

The development of this new Compliance Program Guidance for Clinical Laboratories has been enhanced